## In the Claims:

The current status of all claims is listed below and supersedes all previous lists of claims.

Please cancel claims 13, 15-17, and 22-24 without prejudice to their presentation in another application, amend claims 12-14 and 21, and add new claims 29-32 as follows:

- 1-11. (canceled).
- 12. (currently amended) A vaccine composition comprising a peptide sequence comprising the N-terminal portion of the angiotensin-II type-1 receptor defined by the sequence MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG (SEQ ID NO:1), in which the peptide is conjugated to a carrier protein or a fragment thereof.
- 13. (canceled).
- 14. (currently amended) A method of treating cancer <u>comprising comprising:</u> administering to a subject in need thereof a therapeutically effective amount of a monoclonal antibody, or a fragment thereof, that binds to a peptide;

wherein the peptide comprises an N-terminal portion of an angiotensin-II type-1 receptor comprising the sequence MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG (SEQ ID NO:1), a conservative mutant thereof, or an active fragment thereof comprising at least five amino acid residues.

- 15-17. (canceled).
- 18. (previously presented) The method of claim 14 wherein the monoclonal antibody is humanized.

- 19. (previously presented) The method of claim 14 wherein the monoclonal antibody is 6313/G2 produced by the hybridoma cell line designated by accession number 93072117.
- 20. (previously presented) The method of claim 14 wherein the cancer is prostate cancer or breast cancer.
- 21. (currently amended) A method of treating a disease or condition associated with vascular smooth muscle cell proliferation comprising comprising:

administering to a subject in need thereof a therapeutically effective amount of a monoclonal antibody, or a fragment thereof, that binds to a peptide;

wherein the peptide comprises an N-terminal portion of an angiotensin-II type-1 receptor comprising the sequence MILNSSTEDG IKRIQDDCPK AGRHNYIFVM IPTLYSIIFV VGIFG (SEQ ID NO:1), a conservative mutant thereof, or an active fragment thereof comprising at least five amino acid residues.

- 22-24. (canceled).
- 25. (previously presented) The method of claim 21 wherein the monoclonal antibody is humanized.
- 26. (previously presented) The method of claim 21 wherein the monoclonal antibody is 6313/G2 produced by the hybridoma cell line designated by accession number 93072117.
- 27. (previously presented) The method of claim 21 wherein the disease or condition is atherosclerosis.
- 28. (previously presented) The composition of claim 12 further comprising an adjuvant.

- 29. (new) A composition comprising a peptide comprising up to 45 amino acids, and comprising the sequence EDGIKRIQDD (SEQ ID NO:2).
- 30. (new) The composition of claim 29 in which the peptide is conjugated to a carrier protein.
- 31. (new) The method of claim 14 wherein the antibody fragment is a Fab, F(ab')<sub>2</sub>, Fv, or scFv fragment.
- 32. (new) The method of claim 21 wherein the antibody fragment is a Fab, F(ab')<sub>2</sub>, Fv, or scFv fragment.